**Amendments to the Claims:** 

This listing of claims will replace all prior versions, and listings, of claims in the

application:

**Listing of Claims:** 

Claim 1 (currently amended): A method for immobilizing biomolecules, which

method comprises contacting a solution containing a biomolecule or biomolecules

provided with at last least one tag with an immobilization substrate which has (i)

binding sites for the biomolecule tag or tags, and (ii) activated reactive groups which

are capable of forming covalent bonds with the biomolecule or biomolecules.

Claim 2 (original): The method according to claim 1, comprising the steps of:

a first step wherein the reactive groups of the immobilization substrate which

are capable of forming a covalent bond with the biomolecule or biomolecules to be

immobilized are activated;

a second step wherein a solution containing the biomolecule or biomolecules

to be immobilized is reacted with the immobilization substrate following the first step,

and

wherein, in the second step, the biomolecule or biomolecules are immobilized on the

immobilization substrate through interaction between the tag or tags and tag-binding

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sites of the immobilization substrate and covalent bonds formed between the reactive

groups and the biomolecule or biomolecules.

Claim 3 (original): The method according to claim 2, wherein the reactive groups are

carboxyl groups, and in the second step, an amine coupling is formed between the

carboxyl groups and an amino group on the biomolecule to be immobilized.

Claim 4 (previously presented): The method according to claim 2, wherein the tag is a

histidine tag, and in the second step, an interaction is effected between the histidine

tag and the immobilization substrate.

Claim 5 (original): The method according to claim 4, wherein, in the second step, an

interaction is effected between the histidine tag and the immobilization substrate

through a complex.

Claim 6 (original): The method according to claim 5, wherein, in the second step, an

interaction is effected between the histidine tag and the immobilization substrate

through a metal ion chelate.

Claim 7 (original): The method according to claim 6, wherein, in the second step, an

interaction is effected between the histidine tag and the immobilization substrate

through Ni<sup>2+</sup> nitrilotriacetic acid (Ni-NTA).

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Claim 8 (original): The method according to claim 6, wherein, in the second step, an

interaction is effected between the histidine tag and the immobilization substrate

through Ni<sup>2+</sup> iminodiacetic acid (Ni-IDA).

Claim 9 (previously presented): The method according to claim 1, wherein the tag-

binding site of the immobilization substrate is an antibody to the tag.

Claim 10 (original): The method according to claim 9, wherein the tag is a histidine

tag, the antibody is an anti-histidine antibody and, in the second step, an intereaction

is effected between the histidine tag and the immobilization substrate through an anti-

histidine antibody.

Claim 11 (previously presented): The method according to claim 1, wherein the tag is

an inherent part of a native biomolecule.

Claim 12 (previously presented): The method according to claim 1, wherein the

biomolecule is a protein.

Claim 13 (previously presented): A method for determining biomolecule-low

molecular weight compound affinity and/or kinetics comprising:

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a step for reacting a sample containing a low molecular weight compound or

compounds to be determined with an immobilization substrate to which a biomolecule

or biomolecules have been immobilized using the method for immobilizing

biomolecules as defined in claim 1, and

a step for determining the affinity and/or kinetics of the low molecular weight

compound or compounds contained in the sample for the biomolecule or

biomolecules immobilized on the immobilization substrate.

Claim 14 (original): The method according to claim 13, wherein the affinity and/or

kinetics of a biomolecule and a low molecular weight compound is determined using

the principle of surface plasmon resonance (SPR) in the step for determining affinity

and/or kinetics.

Claim 15 (previously presented): The method according to claim 13, wherein the

biomolecule is a protein.

Claim 16 (withdrawn): A method for determining protein-protein affinity and/or

kinetics comprising:

a step for reacting a sample containing a protein or proteins to be determined

with an immobilization substrate which has a protein or proteins immobilized thereon

using the method for immobilizing biomolecules as defined in claim 1, and

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a step for determining the affinity and/or kinetics of the protein or proteins

contained in the sample for the protein or proteins immobilized on the immobilization

substrate.

Claim 17 (withdrawn): The method according to claim 16, wherein the affinity and/or

kinetics of a protein in the sample for an immobilized protein is determined using the

principle of surface plasmon resonance (SPR) in the step for determining the affinity

and/or kinetics.

Claim 18 (previously presented): An immobilization substrate comprising at least one

immobilized biomolecule, wherein the biomolecule or biomolecules have been

immobilized by the method defined in claim 1.

Claim 19 (original): The immobilization substrate of claim 18, which comprises:

a substrate, and

polysaccharide chains arranged on the substrate, into which are introduced

reactive groups capable of forming covalent bonds with a biomolecule or

biomolecules to be immobilized thereon,

wherein the biomolecule or biomolecules interact with the polysaccharide chain

through a chelate and form covalent bonds with the reactive groups.

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Claim 20 (previously presented): The method according to claim 18, wherein the biomolecule is a protein.